

IN THE SPECIFICATION

Please replace the paragraph beginning on page 54, line 18, with the following:

1,3-Dimethoxy-5-iodo-2-isopropoxybenzene (2.25 g) was treated in the same manner as described in Preparation Example [[27]] 29 to give the title compound.

Yield: 1.23 g (74%).

Please replace the paragraph beginning on page 70, line 19, with the following:

1-[[2-(3,4,5-Trimethoxyphenyl)pyridin-4-yl]methyl-4-piperidone (1.40 g) and 3,4-methylenedioxyaniline (646 mg) were treated in the same manner as described in Preparation Example [[29]] 37 to give the title compound.

Yield: 810 mg (43%).

Please replace the paragraph beginning on page 87, line 10, with the following:

4-Anilino-1-[[2-(3,4,5-trimethoxyphenyl)pyridin-4-yl]methyl]piperidine (1.64 g) and 4-chloromethyl-2-(3,4,5-trimethoxyphenyl)pyridine (1.64 g) were reacted in the same manner as described in ~~Preparation~~ Example 9. The title compound was obtained after converting the product to a trihydrochloride.

Yield: 635 mg (20%).

Please replace the paragraph beginning on page 126, line 11, with the following:

4-(*m*-Anisidino)-1-(tert-butoxycarbonyl)piperidine (613 mg) and 3-(3,4,5-trimethoxyphenyl)benzyl chloride (586 mg) was treated in the same manner as described in ~~Preparation~~ Example 9 to give light yellow amorphous of the title compound.

Yield: 1.01 g (90%).

Please replace the paragraph beginning on page 157, line 5, with the following:

1-(tert-Butoxycarbonyl)-4-piperidone (5.00 g) and 4-bromoaniline (4.11 g) was treated in the same manner as described in Preparation Example 37 to give white crystalline powder of the title compound.

Yield: 3.09 g (36%).

Please replace the paragraph beginning on page 194, line 10, with the following:

These compounds were obtained by the condensation of amines obtained in Preparation Examples 96, 205, 207, 209, 211, 213, 215, 217, 219, 221, 223 and 225 with chloride derivatives obtained in Preparation Examples 3, ~~93, 94, 95, 96, 97, 98, 99, 100, 101,~~ ~~102~~ 193, 194, 195, 196, 197, 198, 199, 200, 201, 202 and ~~103~~ 203. Free bases obtained were then converted to the corresponding hydrochlorides. Yields and NMR data of their free bases are listed below.

Please replace the paragraph beginning on page 204, line 6 under Example 204, with the following:

Synthesis of ~~4-[N-(4-methoxyphenyl)-N-[[2-(3,4,5-trimethoxyphenyl)pyridin-4-yl]methyl]amino]-1-[[2-(3-methoxyphenyl)pyridin-4-yl]methyl]piperidine trihydrochloride~~
4-[N-(4-methoxyphenyl)-N-[[3-(3,4,5-trimethoxyphenyl)pyridin-5-yl]methyl]amino]-1-[[2-(3-methoxyphenyl)pyridin-4-yl]methyl]piperidine trihydrochloride:

Please replace the paragraph beginning on page 204, line 20 under Example 205, with the following:

Synthesis of ~~4-[N-(4-methoxyphenyl)-N-[[2-(3,4,5-trimethoxyphenyl)pyridin-4-yl]methyl]amino]-1-[[2-(3,4-dimethoxyphenyl)pyridin-4-yl]methyl]piperidine~~

~~trihydrochloride~~ 4-[N-(4-methoxyphenyl)-N-[[3-(3,4,5-trimethoxyphenyl)pyridin-5-yl]methyl]amino]-1-[[2-(3,4-dimethoxyphenyl)pyridin-4-yl]methyl]piperidine trihydrochloride:

Please replace the paragraph beginning on page 207, line 17, with the following:

~~4-[N-(4-Methoxyphenyl)-N-[[3-(3,4,5-trimethoxyphenyl)pyridin-5-yl]methyl]amino]piperidine dihydrochloride~~ 4-[N-(4-Methoxyphenyl)-N-[[3-(3,4,5-trimethoxyphenyl)pyridin-5-yl]methyl]amino]piperidine dihydrochloride (121 mg, obtained in the Preparation Example 143) and ~~4-chloromethyl-2-(4-methoxyphenyl)pyridine~~ 4-chloromethyl-2-(3-methoxyphenyl)pyridine (55 mg, obtained in the Preparation Example 195) were condensed in the same manner described in the Example 9 to give the title compound.

Yield: 71 mg (44%).